From: Coughlin, Daniel M. [dan.coughlin@walgreens.com]

**Sent**: 8/3/2010 1:05:17 PM

To: Ranick, Marcella F. [marcie.ranick@walgreens.com]

CC: Martin, Barbara A. [barb.martin@walgreens.com]; Murray Jr, Denman D. [denman.murray@walgreens.com]; Peters,

Gary A. [gary.peters@walgreens.com]; Pinon, Dwayne A. [dwayne.pinon@walgreens.com]; Svihra, Edward J.

[ed.svihra@walgreens.com]; Jacobs, Robert C. [robbie.jacobs@walgreens.com]; Gorman, Timothy J. [tim.gorman@walgreens.com]; Khanna, Rakesh [rakesh.khanna@walgreens.com]; Stahmann, Eric A.

[eric.stahmann@walgreens.com]

Subject: Re: Suspicious Controlled Drug Orders

Attachments: image3.Gif; image4.Gif; image1.Gif; image2.Gif; image0.Gif

Travelling and unable to view. Some brief question though:

- 1. I recall the old paper report as being inches thick. This was replaced by same data on disc and eventually electronic tr ansmission. We were instructed in 1985 not to review or contact anyone on the data. Who from your group has been reviewing the data collected for the past twenty-five years?
- 2. Doesn't the new program stop the release of suspicious quantity? Wouldn't this give your team more time or ability for whatever review would be performed? At present is anyone doing any review on what would be considered suspicious quantities that are physically ordered and are releasing to stores?

---- Original Message -----

From: Marcie Ranick

Sent: 08/03/2010 12:38 PM CDT

To: Dan Coughlin

Cc: Barb Martin; Denman Murray; Gary Peters; Dwayne Pinon; Ed Svihra; Robbie Jacobs;

Tim Gorman; Rakesh Khanna; Eric Stahmann

Subject: Re: Suspicious Controlled Drug Orders

Dan - in addition to the issues sent previously, the screens below show a discrepancy in the data which is compiled in the new ADR4 -

Threshold Violations program compared to the existing Suspicious Controlled Drug "Trigger" data, which is currently on M obius and is sent to the DEA on a monthly basis. From these screens, you can see that the new Threshold Violations rep ort does not match the data on Mobius Suspicious Controlled Drug Orders.

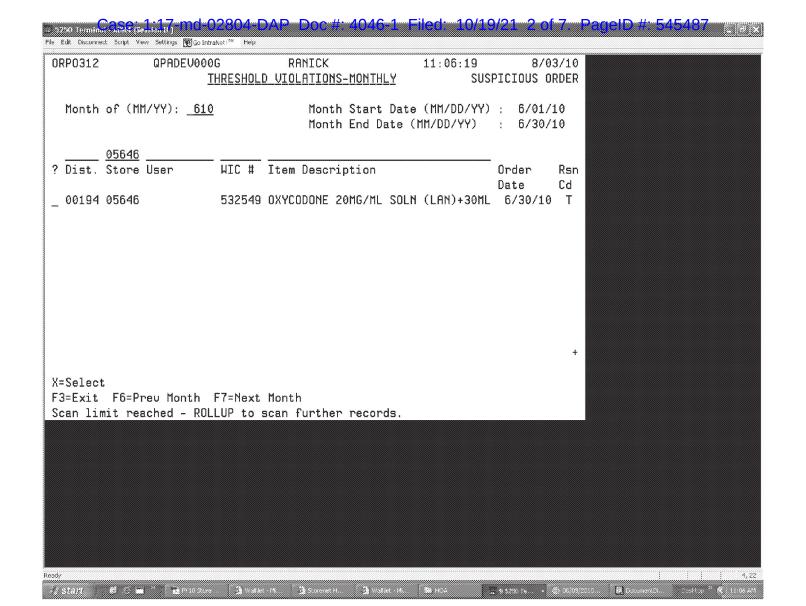
While it is understood that the new calculations will offer different results, it is apparent that the Mobius report will continue to list potential suspicious activity that is not addressed in the new ADR4 system. From this perspective, we believe that the new system should be tested further and enhanced to provide broader coverage of controlled substance activity.

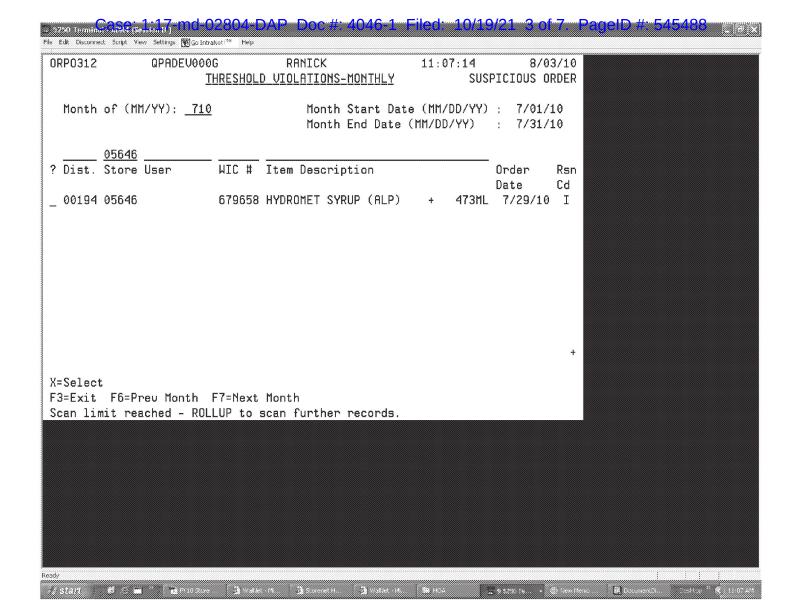
Although we will continue to explore the issues and results of the new system, we are not equipped to handle the 389+ pages of ADR4 data which are compiled nationwide each week. Is there a resource available in your department for data an alysis?

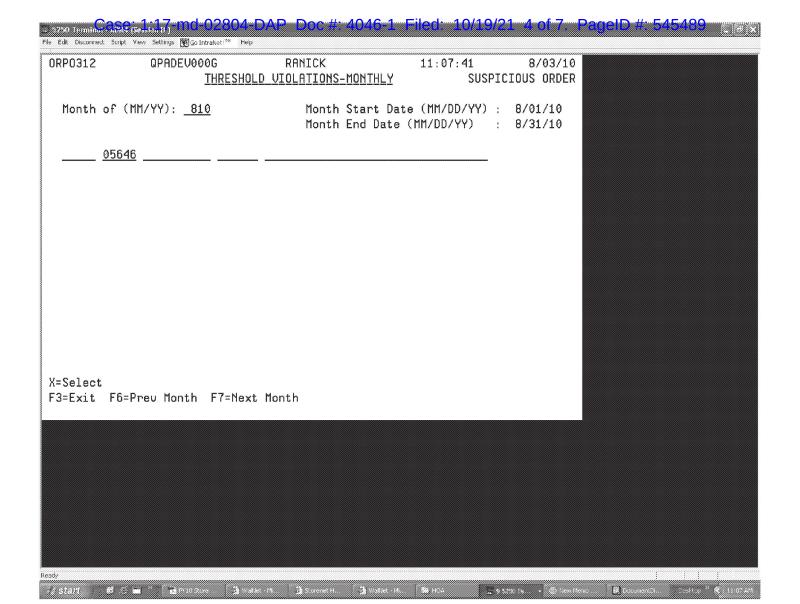
Please let me know if you have any questions, or would like to discuss. I would be happy to set up a meeting to discuss full rollout, a date for the discontinuation of the monthly Mobius CD-disc distribution, and subsequent monitoring and reporting from the new system.

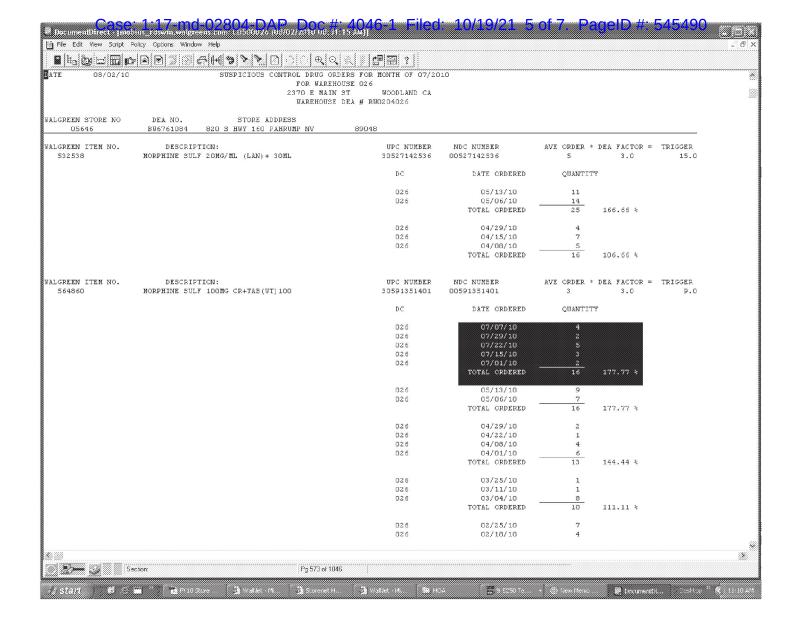
Confidential - this report and its content is intended for restricted Walgreens internal distribution only.

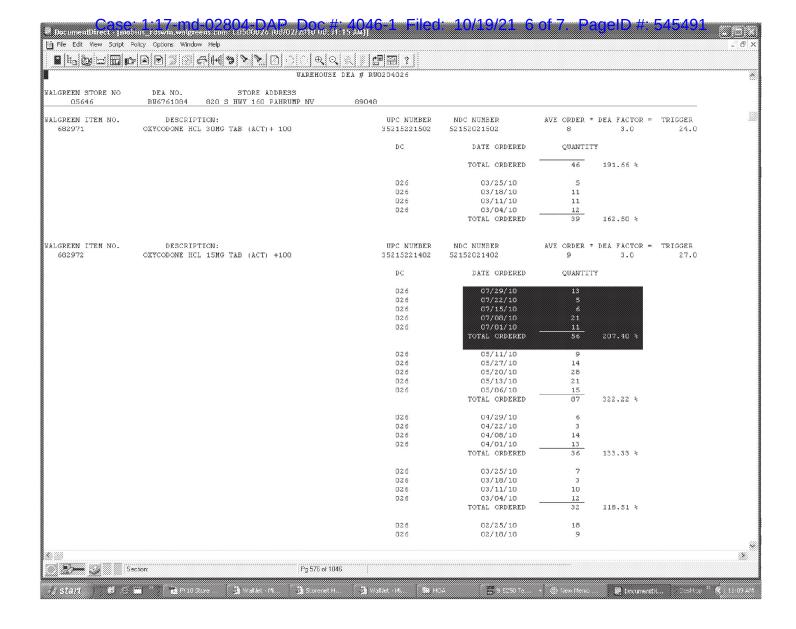
PLAINTIFFS TRIAL EXHIBIT P-00058\_00001











CC

Marcie Ranick, RPH Manager, Pharmacy Loss Prevention 847-964-4193 Fax 847-964-4966

## Dan Coughlin/LOG/Walgreens

07/27/2010 12:51 PM

To Marcie Ranick/Corp/Walgreens@Walgreens, Barb Martin/Corp/Walgreens@Walgreens
Dwayne Pinon/Corp/Walgreens@Walgreens, Denman Murray/Corp/Walgreens@Walgreens, Robbie Jacobs/Corp/Walgreens@Walgreens, Tim Gorman/Corp/Walgreens@Walgreens, Ed Svihra/Corp/Walgreens@Walgreens

WAGMDL00660336

The DEA asked us in 2009 to stop what was considered suspicious drug shipments to any of our stores. This would repla ce the current program where we would send the DEA diversion offices records on disks of what these same transactions were. I believe we have been testing the program that would stop orders considered suspicious in Las Vegas for some time.

Based on the results of Las Vegas, are we ready to roll out chain wide to meet the DEA requirements? From our division, we support the program if tests results have yielded the control of orders while not missing customers' script fills. Our concern is that the DEA could walk into a DC and discuss our inability to have the process in place after their request was made last year.

Any issues that would prevent presentation to the Market Pharmacy Directors and Executive group for pharmacies so they are aware of the changes before chain wide roll out?